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(54) Controlled release formulations of ranitidine

(57) In oral formulations in form of coated tablets or capsules of the therapeutically active drug Ranitidine Hydrochloride, designed or intended to provide release of the active drug in a controlled manner over longer periods and containing pharmacologically acceptable dose of Ranitidine Hydrochloride blended with specified polymers, the quantity of the specified polymer is within the ratio of 0.2: 1.2 with the quantity of the active drug, Ranitidine Hydrochloride.



EUROPEAN SEARCH REPORT

Application Number EP 96 10 0278

Category	Citation of document with i of relevant pass	ndication, where appropriate, ages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (InlCI.6)
Х	US 5 407 687 A (COI	FFIN MARK D ET AL)	1-3,6,7, 9-11	A61K9/52
Y	* abstract * * column 2, line 49-59 *		4,5,8, 12-14	A61K31/34
	* column 3, line 46 * column 3, line 34 * examples 1,3 *			
х	WO 92 10173 A (SMIT * page 2, line 7 *	THKLINE BEECHAM CORP)	1-3	
Υ	* page 3, line 28 - * page 7, line 13 - * page 8, line 6 * * claim 4 *	line 30 * line 23 *	13	
Y	EP 0 446 753 A (VEC * page 3, line 20 - * page 4, line 20 - * page 5, line 16 *	- line 48 * - line 57 *	8,12,14	
	* claim 2 *		_	TECHNICAL FIELDS SEARCHED (Int.Cl.6)
Y	* page 3, line 22 -		D) 4,5	A61K
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.	The present search report has	been drawn up for all claims		
,	Place of search	Date of completion of the sear		Examiner Contago D
	THE HAGUE	9 December 19		Gaetana, R
X : parti Y : parti	ATEGORY OF CITED DOCUMENTS cularly relevant if taken alone cularly relevant if combined with anot ment of the same category	E : earlier paix after the fili her D : document	rinoiple underlying the is ant document, but publis ng date cited in the application cited for other reasons	